WHAT IS CLAIMED IS:

1. A method for vaccinating a human against a human immunodeficiency virus comprising the steps of:

selecting an immunogen competent to induce a protective immune response in said human against said human immunodeficiency virus, and

administering to said human an effective amount of said immunogen sufficient to induce a sustained cell mediated immune response against said human immunodeficiency virus.

- 2. The method of claim 1 wherein said Immunogen is an attenuated form of human immunodeficiency virus.
- 3. The method of claim 2 wherein said immunogen has been attenuated by removing all or part of the <u>nef</u> gene from the nucleic acid of said human immunodeficiency virus.
- 4. The method of claim 1 wherein said immunogen is a subunit of said human immunodeficiency virus.
- 5. The method of claim 4 wherein said immunogen is a gp120 subunit of said human immunodeficiency virus.
- a gp160 subunit of said human immunodeficiency virus.
- 7. The method of claim 1 wherein said immunogen is an inactivated human immunodeficiency virus.
 - 8. The method of claim 7 wherein said immunogen has been inactivated by removing a sufficient portion of its genetic material so as to render it incapable of replicating.
 - 9. The method of claim 8 wherein the genetic material removed from said human immunodeficiency virus is a portion of a gene coding for a gag nucleocapsid protein.

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- 10. The method of claim 7 wherein said human immunodeficiency virus has been inactivated by exposure to a solution of betapropiolactone.
- 11. The method of claim 1 wherein said immunogen is an infectious form of human immunodeficiency virus administered in a subinfectious amount.
- 12. The method of claim 1 wherein the effective

 10 amount of immunogen administered contains between 100

 attograms and 20 milligrams of p24 gag antigen.
 - 13. The method of claim 2 wherein the effective amount of immunogen administered contains between 10 and 500 femtograms of p24 gag antigen.
 - 14. The method of claim 11 wherein the effective amount of immunogen administered contains between 100 attograms and 500 femtograms of p24 gag antigen.
 - 15. The method of claim 1 wherein a cell mediated response is determined to be present using a T-Cell proliferation assay if the uptake of thymidine by antigenstimulated cells is at least four-fold above background.
 - response is determined to be present using an IL-2 assay if the production of IL-2 by antigen-stimulated cells is at least four-fold above background.
 - 17. A method for vaccinating a human against a human immunodeficiency virus comprising the steps of:
 selecting an immunogen competent to induce a protective immune response in said human against said human immunodeficiency virus, and

administering an effective amount of said immunogen to said human sufficient to induce a cell mediated

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18. A method for vaccinating a human against a mammalian retrovirus comprising the steps of:

selecting an immunogen competent to induce a protective immune response in said mammal against said retrovirus, and

administering an effective amount of said immunogen to said mammal sufficient to induce a cell mediated immune response against said retrovirus but below the level necessary to induce a humoral response.

- 19. The method of claim 18 wherein said retrovirus is a simian immunodeficiency virus.
- 20. The method of claim 18 wherein said mammal is a human.
- 21. The method of claim 20 wherein said retrovirus is HTLV-I.
- 22. The method of claim 20 wherein said retrovirus is HTLV-II.
- 23. The method of claim 20 wherein said retrovirus is foamy virus.
- 24. A vaccine comprising a therapeutically effective dose of an immunogen capable of eliciting a cell-mediated immune response in a human protective against infection by a human immunodeficiency virus.
- 25. A vaccine comprising a dose of immunogen
 35 capable of eliciting a cell-mediated response in a human as measured by a T-cell proliferation assay.

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